

# Medicines and Healthcare products Regulatory Agency

## MANUFACTURER'S AUTHORISATION

<b>1: Authorisation Number</b>	UK MIA(IMP) 30013
<b>2: Name of authorisation holder</b>	PHARMARON BIOLOGICS (UK) LTD
<b>3: Address(es) of manufacturing site(s)</b>	PHARMARON BIOLOGICS (UK) LTD, NATIONAL BIOMANUFACTURING CENTRE, ESTUARY BANKS, ESTUARY COMMERCE PARK, SPEKE ROAD, LIVERPOOL, L24 8RB, UNITED KINGDOM
<b>4: Legally registered address of authorisation holder</b>	PHARMARON BIOLOGICS (UK) LTD, NATIONAL BIOMANUFACTURING CENTRE, ESTUARY BANKS, ESTUARY COMMERCE PARK, SPEKE ROAD, LIVERPOOL, L24 8RB, UNITED KINGDOM
<b>5: Scope of authorisation and dosage forms</b>	ANNEX 1 and/ or ANNEX 2
<b>6: Legal Basis of authorisation</b>	Part 6 of The Medicines for Human Use (Clinical Trials) Regulations 2004 [SI 2004/1031]
<b>7: Name of responsible officer of the competent authority of the member state granting the manufacturing authorisation</b>	Confidential
<b>8: Authorisation Date</b>	03/03/2022
<b>9: Annexes attached</b>	Annex 1 and/or Annex 2

### SCOPE OF AUTHORISATION

#### Annex 2

Name and address of the site:

**PHARMARON BIOLOGICS (UK) LTD, NATIONAL BIOMANUFACTURING CENTRE, ESTUARY BANKS, ESTUARY COMMERCE PARK, SPEKE ROAD, LIVERPOOL, L24 8RB, UNITED KINGDOM**

Human Investigational Medicinal Products
Authorised Operations
MANUFACTURING OPERATIONS (according to part 1) IMPORTATION OF MEDICINAL PRODUCTS (according to part 2)
<b>Part 1 - MANUFACTURING OPERATIONS</b> <b>[ 1.1 ] Sterile Investigational Medicinal Products</b> [ 1.1.3 ] Batch certification <b>[ 1.2 ] Non-sterile investigational medicinal products</b> [ 1.2.2 ] Batch certification <b>[ 1.3 ] Biological investigational medicinal products</b>

- [ 1.3.1 ] Biological medicinal products
    - [ 1.3.1.2 ] Immunological products
    - [ 1.3.1.3 ] Cell therapy products
    - [ 1.3.1.4 ] Gene therapy products
    - [ 1.3.1.5 ] Biotechnology products
    - [ 1.3.1.6 ] Human or animal extracted products
    - [ 1.3.1.8 ] Other biological medicinal products
- Biological Active Starting Materials

**[ 1.6 ] Quality control testing**

- [ 1.6.2 ] Microbiological: non-sterility
- [ 1.6.3 ] Chemical/Physical
- [ 1.6.4 ] Biological

Part 2 - IMPORTATION OF MEDICINAL PRODUCTS

**[ 2.2 ] Batch certification of imported medicinal products**

- [ 2.2.1 ] Sterile Products
    - [ 2.2.1.1 ] Aseptically prepared
    - [ 2.2.1.2 ] Terminally sterilised
  - [ 2.2.2 ] Non-sterile products
  - [ 2.2.3 ] Biological medicinal products
    - [ 2.2.3.2 ] Immunological products
    - [ 2.2.3.3 ] Cell therapy products
    - [ 2.2.3.4 ] Gene therapy products
    - [ 2.2.3.5 ] Biotechnology products
    - [ 2.2.3.6 ] Human or animal extracted products
    - [ 2.2.3.8 ] Other biological medicinal products
- Biological Active Starting Materials